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(54) Title: CLEANING COMPOSITION AND CLEANING OF VEHICLES

(57) Abstract: Vehicle cleaning compositions containing one or more selected components with complexing properties, the use thereof for removing firmly adhering metal dust-containing and/or abraded metal-containing residues and for cleaning wheel rims, the production of corresponding vehicle cleaning solutions and a method for cleaning vehicles which is particularly protective of the surface.

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CLEANING COMPOSITION AND CLEANING OF VEHICLES

The invention relates to vehicle cleaning compositions containing one or more selected components with complexing properties, to the use thereof for removing firmly adhering metal dust-containing and/or abraded metal-containing residues and for cleaning wheel rims, to the production of
5 corresponding vehicle cleaning solutions and to a method of cleaning vehicles which is particularly protective of the surface.

Various things should be noted when cleaning vehicle surfaces. On the one hand, the surfaces may consist of a wide variety of materials. These surfaces may be produced, for example, from glass, rubber, coated
10 surfaces, steel and aluminum, plastics materials, composites, thermoplastic polymer fabrics or from thermoplastic polymer fibers, plastic lenses or a large number of glass or metal composites.

However, many different types of soiling may also occur. This soiling may originate, for example, from fuels, lubricants, hydraulic and
15 other operating fluids, dirt, components of the vehicle exhaust, residues of previously used detergents, waxes and the like. Firmly adhering residues containing metal dust and/or abraded metal are particularly difficult to remove.

Soiled vehicle surfaces have been cleaned for years using a large
20 number of compositions and methods. These compositions may be simple solutions of organic dishwashing detergents or conventional multi-purpose detergents. When cleaning vehicles in trade or industry, for example during the semiautomatic and fully automatic washing of cars, a large number of detergents have been used for cleaning systems, the cleaning
25 systems often involving a prewash cycle or precleaning cycle, a cleaning cycle and then a combination of one or more operations involving wax, detergent, rust-proofing agent, mechanical drying and the like. This type of vehicle cleaning may be the commercial cleaning of vehicles during which

the vehicles are washed by the owner himself or by employees. The cleaning stations may be, for example, stations which are operated by car rental firms, car dealers, the operators of car fleets, in bus and railway depots, in aircraft servicing buildings and the like.

- 5 One category of generally available detergents contains a number of anionic surfactants which are used together with compatible nonionic surfactants, complexing agents, waxes and other components.

The prior art vehicle detergent is suitable for removing most residues.

- 10 In the vehicle industry, however, there are combinations of dirt which contain, for example, grease, carbon black and abraded metal which, after thorough cleaning with detergents which have a high surfactant content and readily dissolve grease, leave a thin, firmly adhering dust film consisting of abraded metal, in particular from brake linings of cars,
15 railways vehicles or the like.

In the past, this dust could only be cleaned with highly alkaline or strongly acidic detergents, while employing, for example, high pressure or brushes.

- However, the use of these detergents adversely affects the surfaces
20 to be cleaned. Strong alkaline solutions and acids may markedly attack the coating on surfaces, for example on car wheel rims and bodies, and the coating on railway carriages. The coated surfaces of car wheel rims or railway carriages or other vehicles which have been cleaned several times in this way soon have a matt appearance and therefore lose the
25 appearance of an as-new vehicle.

It is therefore desirable to have further suitable alternatives for vehicle cleaning, in particular with respect to improved material compatibility and ability to remove difficult-to-remove residues such as metal dust and abraded metal.

- 30 The object of the present invention was accordingly to provide

vehicle cleaning compositions which have good material compatibility and are capable of removing difficult-to-remove residues such as metal dust and abraded metal.

This object is achieved by a vehicle cleaning composition containing
5 one or more components with complexing properties selected from the groups comprising polyamino and polycarboxylic acids and the salts thereof.

Suitable polycarboxylic acids include, for example, polyacrylic acids and copolymers of maleic acid anhydride and acrylic acid and the sodium
10 salts of these polymeric acids. Conventional commercial products include Sokalan® CP 5 and PA 30 from BASF, Alcosperse® 175 and 177 from Alco, LMW® 45 N and SPO2 ND from Norsohaas. Suitable natural polymers include, for example, oxidized starch (e.g. DE 42 28 786) and polyamino acids such as polyglutamic acid or polyaspartic acid, for
15 example from Cygnus, Bayer, Rohm & Haas, Rhône-Poulenc or SRCHEM.

The aforementioned components with complexing properties are particularly preferably selected from polyaspartic acid and/or copolymers based on aspartic acid.

It is also preferred if the vehicle cleaning composition according to
20 the invention contains gluconic acid as an additional component with complexing properties. The vehicle cleaning compositions according to the invention may also contain a large number of other materials which are useful in the production of agents for cleaning vehicles and for water removal, such as nonionic surfactants, amine oxide surfactants, further
25 complexing agents, acidic materials, basic materials, solvents and a large number of other useful materials such as colorants, perfumes, thickeners, foaming surfactants, etc.

Conventional hardening or solidifying agents such as urea, PEG materials, nonionic surfactants and the like may also be used. It is
30 preferable if the vehicle cleaning composition according to the invention

additionally contains a nonionic surfactant which is particularly preferably selected from the group comprising alkoxylated fatty alcohols which are optionally terminated by end groups and/or alkyl polyglycosides and/or alkoxylated fatty amines.

5 Suitable alkoxylated fatty alcohols include C₈₋₁₈ alkyl polyethylene glycol polypropylene glycol ethers with up to 8 mol ethylene oxide (=EO) and propylene oxide (=PO) units in the molecule respectively. The addition of tallow alcohol ethoxylated with 30 EO groups and the addition of oleyl-cetyl alcohol ethoxylated with 5 EO groups has also had a positive
10 influence on the cleaning result. However, other known nonionic surfactants may also be used, such as C₁₂₋₁₈ alkyl polyethylene glycol polybutylene glycol ethers with up to 8 mol ethylene oxide and butylene oxide units in the molecule respectively, as well as alkyl polyalkylene glycol mixed ethers terminated by end groups. Suitable alkoxylated fatty amines
15 include, for example, C₈₋₁₈ alkyl amines ethoxylated with 8 to 16 EO groups.

A further important nonionic surfactant can comprise an amine oxide. Representative examples of such amine oxides include lauryldimethyl amine oxide, dodecyldimethyl amine oxide,
20 tetradecyldimethyl amine oxide, cetyldimethyl amine oxide, stearyldimethyl amine oxide, dodecyldiethyl amine oxide, bis(2-hydroxypropyl) tetradecyl amine oxide and the like.

The pH of the vehicle cleaning composition according to the invention preferably ranges from 4 to 10, particularly preferably from 5 to 9
25 and more particularly preferably from 6 to 8.

Preferred embodiments of the vehicle cleaning composition according to the invention contain 0.1 to 75 % by weight, particularly preferably 0.5 to 50 % by weight of the aforementioned complexing agents contained according to the invention, based on the total weight of vehicle
30 cleaning composition.

Particularly preferred embodiments of the vehicle cleaning composition according to the invention contain 0.1 to 30 % by weight of polyaspartic acid, more particularly preferably in combination with 0.1 to 50 % by weight of gluconic acid, based on the total weight of vehicle cleaning composition.

The present invention also relates to the use of vehicle cleaning compositions according to the invention for the cleaning of vehicle surfaces, in particular for the removal of metal dust-containing and/or abraded metal-containing residues adhering firmly to vehicle surfaces. The use according to the invention is particularly preferred if the residues consist of iron oxide or at least contain iron oxide fractions.

Surfaces include, in particular, vehicle wheel rims and any other surfaces mentioned in the description of this application, such as coated surfaces and the like.

For the purposes of this description, the term "vehicle" refers to any means of transport, including cars, lorries, sports/utility vehicles, buses, golf buggies, motor cycles, monorail vehicles, diesel engines, long-distance coaches, small single-engine private aircraft, airline jet planes, conventional commercial airline equipment and the like.

With the use according to the invention, the removal of metal dust-containing and/or abraded metal-containing residues adhering firmly to vehicle surfaces is preferably carried out in a separate cleaning stage detached from the conventional cleaning of vehicles, before or after the conventionally performed cleaning of the vehicles to be cleaned.

The present invention also relates to the production of vehicle cleaning solutions by dilution of a vehicle cleaning composition according to the invention with water, so that the ready-to-use cleaning solution contains 0.025 to 100 g/l, preferably up to 50 g/l of at least one of the aforementioned components with complexing properties and, in a particularly preferred embodiment, additionally contains 0.05 to 50 g/l, more

particularly preferably 0.5 to 5 g/l of the aforementioned nonionic surfactant.

The present invention also relates to vehicle cleaning solutions according to the invention produced by mixing a solution containing at least one of the aforementioned components having complexing properties with a solution containing the aforementioned nonionic surfactant, a ready-to-use cleaning solution containing 0.025 to 100 g/l, preferably 50 g/l, of at least one of the aforementioned components with complexing properties and additionally 0.05 to 50 g/l, more particularly preferably 0.5 to 5 g/l, of the aforementioned nonionic surfactant being obtained, optionally after further dilution.

Finally, the present invention also relates to a method for cleaning a vehicle which is particularly protective of the surface and in which a vehicle cleaning solution according to the invention is applied to the surface of the vehicle and is rinsed off with water after the cleaning operation.

The vehicle cleaning compositions according to the invention are preferably used in systems which operate with brushes or by contact with cloth or without contact. Contactless systems usually operate by merely spraying the aqueous systems, followed by rinsing with water which leaves a cleaner vehicle surface containing small amounts or no residues of the cleaning composition or the detergent.

The vehicle cleaning compositions according to the invention are typically applied to a vehicle surface to remove many types of soiling which usually occurs during transportation, during transfer by rail, at airports and on motorways and the like.

The term "contactless cleaning system" denotes a method with which the detergents are brought into direct contact with a vehicle surface, including coated surfaces, surfaces made of thermoplastic laminate, glass surfaces, plastic surfaces or surfaces comprising conventional car trims, in order to remove soil by spraying or washing without a mechanical action

being employed during the removal of dirt. In use, an aqueous cleaning composition has to remain in contact with these surfaces for a relatively short time (shorter than 5 minutes) in order to assist dirt removal. The aqueous systems are typically removed from the vehicle surface by rinsing
5 with water and then with a water removing agent.

The vehicle cleaning compositions according to the invention may generally be applied in liquid, solid, paste or other forms.

The vehicle cleaning compositions according to the invention may contain acidic or basic substances as further ingredients, with which a
10 basic or acidic pH may be neutralized. These basic substances include, for example, amines, sodium hydroxide, sodium silicates and the like. These materials may act as alkaline builders, dirt carriers and as buffers. The preferred silicates may also act as protection for aluminum by reducing the effects of the chemicals according to the invention on exposed aluminum
15 surfaces. The alkaline builders should be present in sufficient large amounts for achieving a substantially neutral pH (for example about 6 to 10, preferably 6 to 9). A large number of typical weak or mild acids may be used to neutralize and solubilize the basic compositions to different pH values. These acids include, for example, acetic acid, hydroxyacetic acid,
20 phosphoric acid, citric acid and other typical acids which are used to produce cleaning compositions.

The vehicle cleaning compositions according to the invention may contain a solvent. The preferred chemistry requires no solvent. Preferred solvents include alcohols, glycols and glycol ethers. These substances
25 generally have an aliphatic fraction containing 2 to 6 carbon atoms. Examples of these substances include ethanol, propanol, isopropanol, butanol, 2-butanol, 2-methyl-2-propanol, butoxydiglycol, ethoxydiglycol, polypropyleneglycol, ethyleneglycol methylether, ethyleneglycol dimethylether, propyleneglycol methylether, dipropyleneglycol-n-butylether,
30 butoxyethanol, phenoxyethanol, methoxypropanol, propyleneglycol, n-

butylether, tripropyleneglycol, n-butylether, propyleneglycol, hexyleneglycol and other similar oxygen-containing solvents.

Examples

5 The following test was carried out in order to discover the effectiveness of different detergent formulations against soiling which is typical of the aforementioned applications:

Some stainless steel panels (5 x 10 cm) were prepared for the test by applying 0.4 to 0.5 g of standard dirt to one side of the test panel and then leaving it to dry for 24 hours at 25 °C. Car brake dust combined with
10 mineral oil dirt (1), on the one hand, and railway brake dust combined with iron oxide dirt (2), on the other hand, were used as standard dirt.

The cleaning test was carried out by immersing the test objects pretreated in this way into the various cleaning solutions at temperatures of 20 °C for 20 minutes in a fully automatic immersion apparatus. The
15 detachment of the covering was determined gravimetrically.

The formulations shown in the following table were converted, by dilution with water to service concentrations, into a cleaning solution of which the cleaning capacity was determined in the test.

Table 1:

20 Cleaning capacity of formulations of different compositions (details in % by weight) with respect to soil conventionally encountered in the engineering industry

Ingredients	Examples/active ingredient content in wt. %						
	Comp. 1	Comp. 2	Comp. 3	Comp. 4	Ex. 1	Ex. 2	Ex. 3
Citric acid	0	20		50		5	
NaOH (50 %)	25	0	25	0	2	0	0
NTA	20	0	0	0	4	0	0
EDTA	0	20	20	0	0	2	0
Gluconic acid	0	0	0	0	10	25	25
Hydroxyethane diphosphonate (Na salt)	2.5	2.5	2.5	2.5	2.5	2.5	2.5
Polyaspartic acid	0	0	0	0	10	20	20
C ₁₂₋₁₈ fatty alcohol 9 EO Butyl ether				16.5			
C ₁₂₋₁₈ fatty alcohol 5 EO + 4 PO	15	15	15		15	15	15
Coconut amine 12 EO	0	0	0	16.5	0	5	5
Alkyl glycoside	2	1	2	0	5	0	0
Remainder to 100 %	Demineralized water						
SW concentration (%)	4	4	4	4	4	4	4
Soil removal							
Test soil (1)	11.3	1.5	11.8	16.4	78	98.5	98.7
Test soil (2)	12.6	2.8	12.6	19.3	89	99.6	99.6

Comp. 1 to 4 = comparison examples

Ex. 1 to 3 = examples according to the invention

The material compatibility was also investigated using the
5 formulations listed in Table 1. On the one hand, the material compatibility was investigated on stainless steel test panels (5 x 10 cm) coated with polyacrylate resin coatings conventional in the vehicle industry. On the other hand, the tests were carried out on aluminum panels (5 x 10 cm).

The DLG (Deutsche Landwirtschafts-Gesellschaft) rotary test was
10 used in all cases. Cleaning solutions containing 2 % by weight of the respective formulation were produced by dilution of the formulation listed in Table 1 with 16 °d water, for carrying out the tests.

The prepared aluminum panels and coated stainless steel test

panels were then cleaned at 25 °C for a contact time of 10 minutes in accordance with the DLG rotary test. They were then rinsed for 2 minutes with water and subsequently dried for 2 minutes. 10 cleaning cycles were carried out in this way.

- 5 The attack on the test panels was then evaluated visually. The results of the evaluation are compiled in Table 2. The evaluation numbers have the following meanings:

- 1 = no attack in comparison with zero (16 °d water)
 2 = slight, acceptable matting of the surface
 10 3 = unacceptable matting of the surface
 4 = clearly visible attack of the surface
 5 = pronounced attack of the surface
 6 = complete destruction of the surface.

Table 2:

- 15 Material compatibility of aqueous cleaning solutions containing 2 % by weight of formulations of different compositions (according to Table 1) by the DLG rotary test relative to aluminum panels and coated stainless steel panels

Material compatibility	Visual evaluation of the examples						
	Comp. 1	Comp. 2	Comp. 3	Comp. 4	Ex. 1	Ex. 2	Ex. 3
Coated stainless steel panels	5	3	5	4	2	2	1
Aluminum panels	6	4	6	5	2	2	2

Comp. 1 to 4 = comparison examples

- 20 Ex. 1 to 3 = examples according to the invention

As shown clearly by the material compatibility tests, aluminum surfaces and coated stainless steel surfaces are much less subject to attack by the formulations according to the invention than by formulations from the prior art.

CLAIMS

1. A vehicle cleaning composition containing one or more components with complexing properties selected from the groups comprising polyamino and polycarboxylic acids and the salts thereof.
- 5 2. A vehicle cleaning composition according to claim 1, characterized in that it contains polyaspartic acid and/or a copolymer based on aspartic acid as a component with complexing properties.
3. A vehicle cleaning composition according to either of claims 1 or 2, characterized in that it contains gluconic acid as an additional component
10 with complexing properties.
4. A vehicle cleaning composition according to any of claims 1 to 3, characterized in that it contains 0.1 to 75 % by weight, particularly preferably 0.5 to 50 % by weight, of the aforementioned components with complexing properties, based on the total weight of vehicle cleaning
15 composition.
5. A vehicle cleaning composition according to any of claims 1 to 4, characterised in that a nonionic surfactant is additionally contained.
6. A vehicle cleaning composition according to claim 5, characterized in that the nonionic surfactant is selected from the group comprising
20 alkoxylated fatty alcohols which are optionally terminated by end groups and/or alkyl polyglycosides and/or alkoxylated fatty amines.
7. A vehicle cleaning composition according to any of claims 1 to 6, characterized in that the pH of the vehicle cleaning composition ranges from 4 to 10, preferably from 5 to 9 and particularly preferably from 6 to 8.
- 25 8. A vehicle cleaning composition according to any of claims 1 to 7; characterized in that the vehicle cleaning composition contains 0.1 to 30 % by weight of polyaspartic acid, based on the total weight of vehicle cleaning composition.
9. A vehicle cleaning composition according to any of claims 2 to 8,
30 characterized in that the vehicle cleaning composition contains 0.1 to 30 %

by weight of polyaspartic acid and 0.1 to 50 % by weight of gluconic acid, based on the total weight of vehicle cleaning composition:

10. A use of vehicle cleaning compositions according to any one of claims 1 to 9 for cleaning vehicle surfaces, in particular for removing metal
5 dust-containing and/or abraded metal-containing residues which adhere firmly to vehicle surfaces.

11. A use according to claim 10, characterized in that the removal of metal dust-containing and/or abraded metal-containing residues adhering firmly to vehicle surfaces is carried out in a separate cleaning stage
10 detached from the conventional cleaning of vehicles, before or after the conventionally performed cleaning of the vehicles to be cleaned.

12. A method of producing vehicle cleaning solutions by dilution of a vehicle cleaning composition according to any one of claims 1 to 9 with water, so that the ready-to-use cleaning solution contains 0.025 to 100 g/l
15 of at least one of the aforementioned components with complexing properties.

13. A method of producing vehicle cleaning solutions by dilution of a vehicle cleaning composition according to any one of claims 1 to 9 with water, so the ready-to-use cleaning solution contains 0.025 to 100 g/l of at
20 least one of the aforementioned components with complexing properties and additionally 0.05 to 50 g/l of the aforementioned nonionic surfactant.

14. A method of producing vehicle cleaning solutions by mixing a solution containing at least one of the aforementioned components having complexing properties with a solution containing the aforementioned
25 nonionic surfactant, a ready-to-use cleaning solution containing 0.025 to 100 g/l of at least one of the aforementioned components with complexing properties and additionally 0.05 to 50 g/l of the aforementioned nonionic surfactant being obtained, optionally after further dilution.

15. A method for cleaning a vehicle which is particularly protective of the
30 surface and in which a vehicle cleaning solution obtainable according to

any one of claims 12 to 14 is applied to the surface of the vehicle and is rinsed off with water after the cleaning operation.

16. A use of a vehicle cleaning composition according to any one of claims 1 to 9 for the cleaning of wheel rims.

INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C11D3/37 C11D11/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 959 125 A (PINGO ERZEUGNISSE WELSCH & KUFFNER) 24 November 1999 (1999-11-24) paragraphs '0002!', '0009!', '0010!; claim 1; example 2	1-16
X	US 5 770 548 A (LESKOWICZ JAMES J ET AL) 23 June 1998 (1998-06-23) column 1, line 12 -column 2, line 13; examples 1,2	1-16
X	WO 95 24456 A (MONSANTO CO) 14 September 1995 (1995-09-14) page 1, line 21 - line 30; claims; examples	1-16

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Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 892 040 A (NIPPON CATALYTIC CHEM IND ;TEIKOKU CHEMICAL IND CO LTD (JP)) 20 January 1999 (1999-01-20) page 2, line 3-11; examples page 3, line 17 - line 58 -----	1-16
X	DE 199 36 179 A (HENKEL ECOLAB & CO OGH) 8 February 2001 (2001-02-08) page 2, line 19 -page 3, line 31; claims; examples 2,3 -----	1-16

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 03/09086

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0959125	A	24-11-1999	DE 29809172 U1 EP 0959125 A1	06-08-1998 24-11-1999
US 5770548	A	23-06-1998	AU 724432 B2 AU 2990797 A BR 9709453 A CA 2255034 A1 EP 0904342 A1 JP 2000510185 T NZ 332912 A WO 9743372 A1 ZA 9704161 A	21-09-2000 05-12-1997 10-08-1999 20-11-1997 31-03-1999 08-08-2000 28-07-2000 20-11-1997 10-12-1997
WO 9524456	A	14-09-1995	US 5443651 A AT 187481 T CA 2162153 A1 DE 69513750 D1 DE 69513750 T2 EP 0698072 A1 JP 8510507 T WO 9524456 A1	22-08-1995 15-12-1999 14-09-1995 13-01-2000 13-07-2000 28-02-1996 05-11-1996 14-09-1995
EP 0892040	A	20-01-1999	JP 11035921 A JP 11302691 A DE 69811786 D1 DE 69811786 T2 EP 0892040 A2 US 6103686 A	09-02-1999 02-11-1999 10-04-2003 23-10-2003 20-01-1999 15-08-2000
DE 19936179	A	08-02-2001	DE 19936179 A1 AU 6565700 A WO 0109275 A1 EP 1200544 A1	08-02-2001 19-02-2001 08-02-2001 02-05-2002